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Form Approved
OMB NO. 0704-0188

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1. AGENCY USE ONLY (Leave blank)	2. REPORT DATE 12/13/00	3. REPORT TYPE AND DATES COVERED Final Report 12/1/97 - 11/30/99
4. TITLE AND SUBTITLE "Revolutionary Smart Self Assembling Crystalline Colloidal Array Materials for Controlling IR Light Propagation"		5. FUNDING NUMBERS Grant #: DAAG-55-98-1-0025
6. AUTHOR(S) Professor Sanford A. Asher		
7. PERFORMING ORGANIZATION NAMES(S) AND ADDRESS(ES) Department of Chemistry University of Pittsburgh 701 Chevron Science Center 219 Parkman Drive Pittsburgh, Pennsylvania 15260		8. PERFORMING ORGANIZATION REPORT NUMBER
9. SPONSORING / MONITORING AGENCY NAME(S) AND ADDRESS(ES) U.S. Army Research Office P.O. Box 12211 Research Triangle Park, NC 27709-2211		10. SPONSORING / MONITORING AGENCY REPORT NUMBER ARO Proposal #: ARO 38355.1 - PH
11. SUPPLEMENTARY NOTES The views, opinions and/or findings contained in this report are those of the author(s) and should not be construed as an official Department of the Army position, policy or decision, unless so designated by other documentation.		
12a. DISTRIBUTION / AVAILABILITY STATEMENT Approved for public release; distribution unlimited.		12b. DISTRIBUTION CODE
13. ABSTRACT (Maximum 200 words) To develop new syntheses for monodisperse polystyrene colloids.		
14. SUBJECT TERMS C		15. NUMBER OF PAGES 20
		16. PRICE CODE
17. SECURITY CLASSIFICATION OR REPORT UNCLASSIFIED	18. SECURITY CLASSIFICATION OF THIS PAGE UNCLASSIFIED	19. SECURITY CLASSIFICATION OF ABSTRACT UNCLASSIFIED
		20. LIMITATION OF ABSTRACT UL

REVOLUTIONARY SMART SELF ASSEMBLING
CRYSTALLINE COLLOIDAL ARRAY MATERIALS FOR
CONTROLLING IR LIGHT PROPAGATION

FINAL PROGRESS REPORT

PROFESSOR SANFORD A. ASHER

DECEMBER 13, 2000

U.S. ARMY RESEARCH OFFICE

GRANT NUMBER DAAG-55-98-1-0025

DEPARTMENT OF CHEMISTRY
UNIVERSITY OF PITTSBURGH
PITTSBURGH, PA 15260

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TECHNICAL PROGRESS REPORT

A. STATEMENT OF PROBLEM

The work here developed new materials and fabrication methodologies to control IR light propagation.

1. We were to develop new syntheses for monodisperse polystyrene, silica, and PMMA colloids
2. Develop synthetic methods to produce colloidal particles containing quantum dots of CdS, CdTe, etc
3. Develop IR diffracting polymerized crystalline colloidal arrays (PCCA)
4. Develop methods to fabricate PCCA which diffract light in defined wavelengths and bandwidths
5. Fabricate PCCA which are thermally and chemically tunable in the 1-10 μm spectral region
6. Demonstrate thermal tuning and electronic control of PCCA diffraction
7. Develop synthesis of highly charged, monodisperse titania dioxide colloids
8. Fabricate and characterize photonic bandgap crystals

B. SUMMARY OF IMPORTANT RESULTS

1. We developed new methods to synthesize highly charged, monodisperse large polystyrene colloidal particles as reported in enclosed publication 1.
2. We developed novel methods to synthesize monodisperse silica colloids in microemulsions as shown in the synthetic scheme of Fig. 1, and the TEM showing examples of these colloidal particles in Fig. 2. We also developed synthetic methods to form silica particles which contain Ag, or Au quantum dots as shown in Figs. 3 and 4. We functionalized these silica composite particles to make them highly charged, such they self assembled into CCA as shown in Figs. 5 and 6. In addition, we developed methods to form polystyrene colloids which contained Ag quantum dots which also self assembled into CCA (Fig. 7 and 8).

We also developed methods using microemulsions to synthesize large silica particles containing CdS quantum dots (Fig. 9).

3. We developed a number methods to produce PCCA which diffract in the IR. In one method we hydrolyzed the acrylamide PCCA to form carboxylates. At neutral pH the charged carboxylates caused the PCCA to swell due to formation of a Donnan potential. Fig. 10 shows the resultant diffraction shift of a PCCA into the IR as the time for hydrolysis increases. This hydrolysis also creates a PCCA, which operates as a pH and ionic strength sensor. Our publication 2 is a careful study of the physics of the pH and ionic strength swelling, and its utility for sensing pH and ionic strength.

We also developed a chemically reversible disulfide cross linking approach (Fig. 11) and a hydrolysable crosslinking approach (Fig. 12 – 15) to swell PCCA into the IR.

4. The methods discussed above allow us to sensitively control the wavelength of diffraction of the PCCA by carefully adjusting the pH or ionic strength or by decreasing the PCCA crosslinking. We developed methods to control the diffraction bandwidth by altering the scattering power of the colloidal particles by changing their size or refractive index. We also attempted to swell colloidal particles already polymerized into the PCCA. Preliminary attempts to swell polystyrene particles by adding styrene to the PCCA solution indicates success. Further work will be necessary to speed up this process in order to make it practical.

5. We demonstrated that we could achieve thermal tuning by utilizing PCCA synthesized by using hydrogels of N-isopropyl acrylamide.

We began working on electronic control of the PCCA diffraction by utilizing a PCCA which contains an 18-crown-6 crown ether group. This molecular recognition agent binds Pb^{2+} , which causes the PCCA to swell and red shifts the diffraction. We attempted electronic control of the PCCA diffraction by utilizing the electrochemical oxidization of lead metal to Pb^{2+} which would then swell the PCCA into the IR. Preliminary data suggests that this method is viable for the control of PCCA diffraction wavelengths.

6. The most challenging part of the research program involved synthesis of highly charged, monodisperse titania spheres. Although the literature reports synthesis of large, relatively monodisperse titania spheres, we found that none of these recipes were reproducible and did not yield monodisperse colloidal particles. We developed a number of new approaches for synthesizing titania spheres of relatively uniform size, but when we tried to add surface charge the titania colloids suffered, what we think, was a Coulomb explosion where the particles disintegrated into their small primary particles. Only just recently, with funding from other grant support, have we finally succeeded in fabricating monodisperse titania colloidal particles which self assemble into CCA that Bragg diffract light. Fig. 16 shows TEM micrographs of these particles.

7. We reported earlier that colloidal arrays made from polystyrene colloids with a lattice spacing that diffracts ~1200 nm light from the fcc 111 planes diffracts essentially all light of all polarizations at wavelengths below 500 nm. Clearly the refractive index contrast, and the fact that it is an fcc crystal will not allow this sample to be a 3-D photonic bandgap crystal. We have been carefully characterizing the crystal structure and diffraction properties and writing up these results for publication. We expect to submit this work by the end of December, 2000.

C. PUBLICATIONS

1. C. Reese, C. Guerrero, J. Weissman, K. Lee and S. A. Asher, "Synthesis of Highly Charged, Monodisperse Polystyrene Colloidal Particles for the Fabrication of Photonic Crystals", *Journal of Colloid and Interface Science*, **232**, 76-80 (2000)
2. K. Lee and S. A. Asher, "Photonic Crystal Chemical Sensors: pH and Ionic Strength", *Journal of American Chemical Society*, **122**, 9534-9537 (2000)

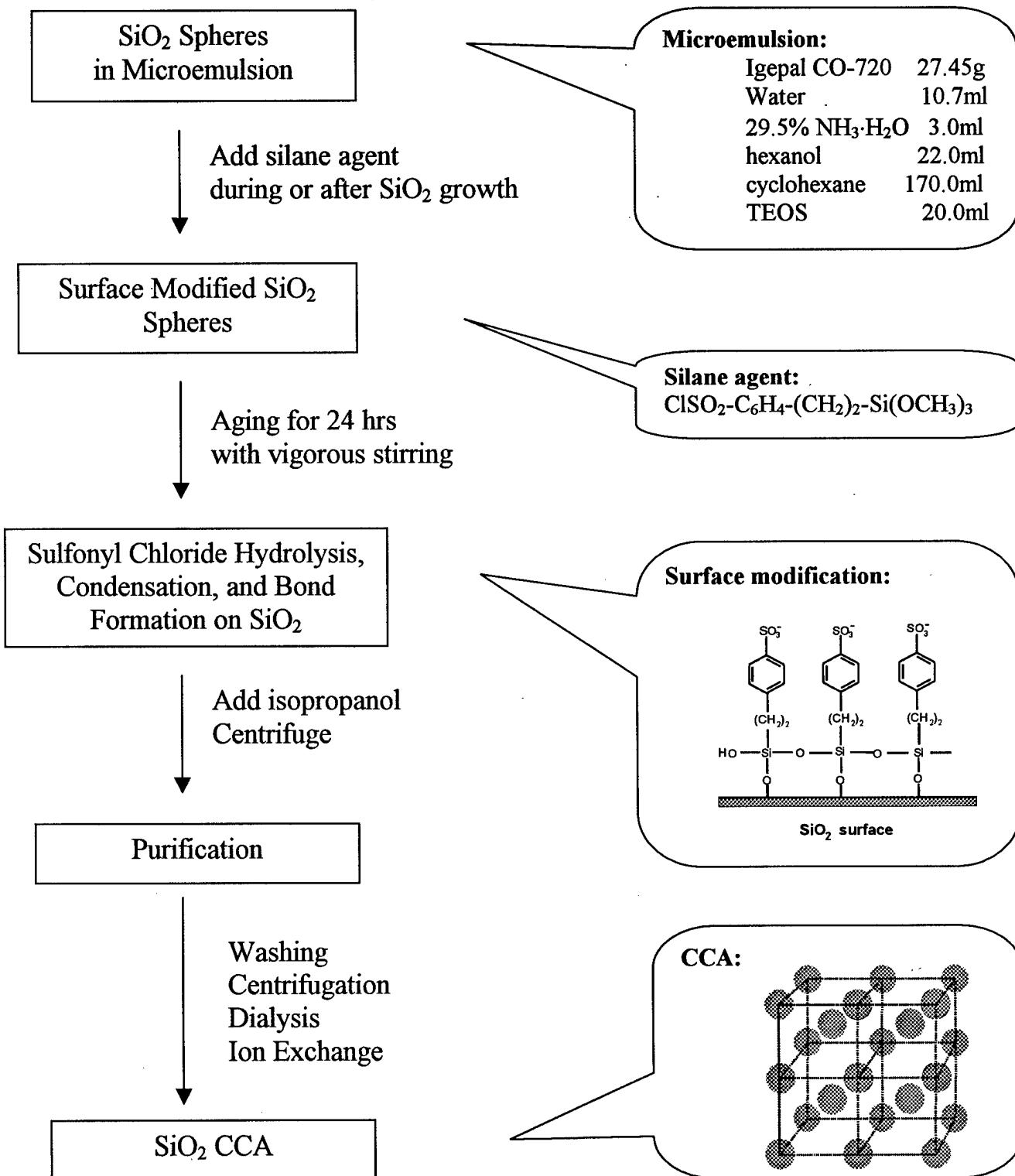
D. LIST OF SCIENTIFIC PERSONNEL EMPLOYED ON PROJECT

Sanford Asher	PI	12%
We Wang	Res. Associate	50%
Igor Lednev	Res. Associate	50%
Jeff Sicher	Machinist	4%
Douglas Hackworth	Graduate Student	75%
Jonathan Keim	Graduate Student	25%
Aleksandr Mikhonin	Graduate Student	25%
Vasil Pajcini	Graduate Student	25% Received PhD
Hua Zhang	Graduate Student	75% Received MS

E. REPORT ON INVENTIONS

None, as yet.

Figure 1. Synthesis of Highly Charged, Monodisperse SiO₂ Particles



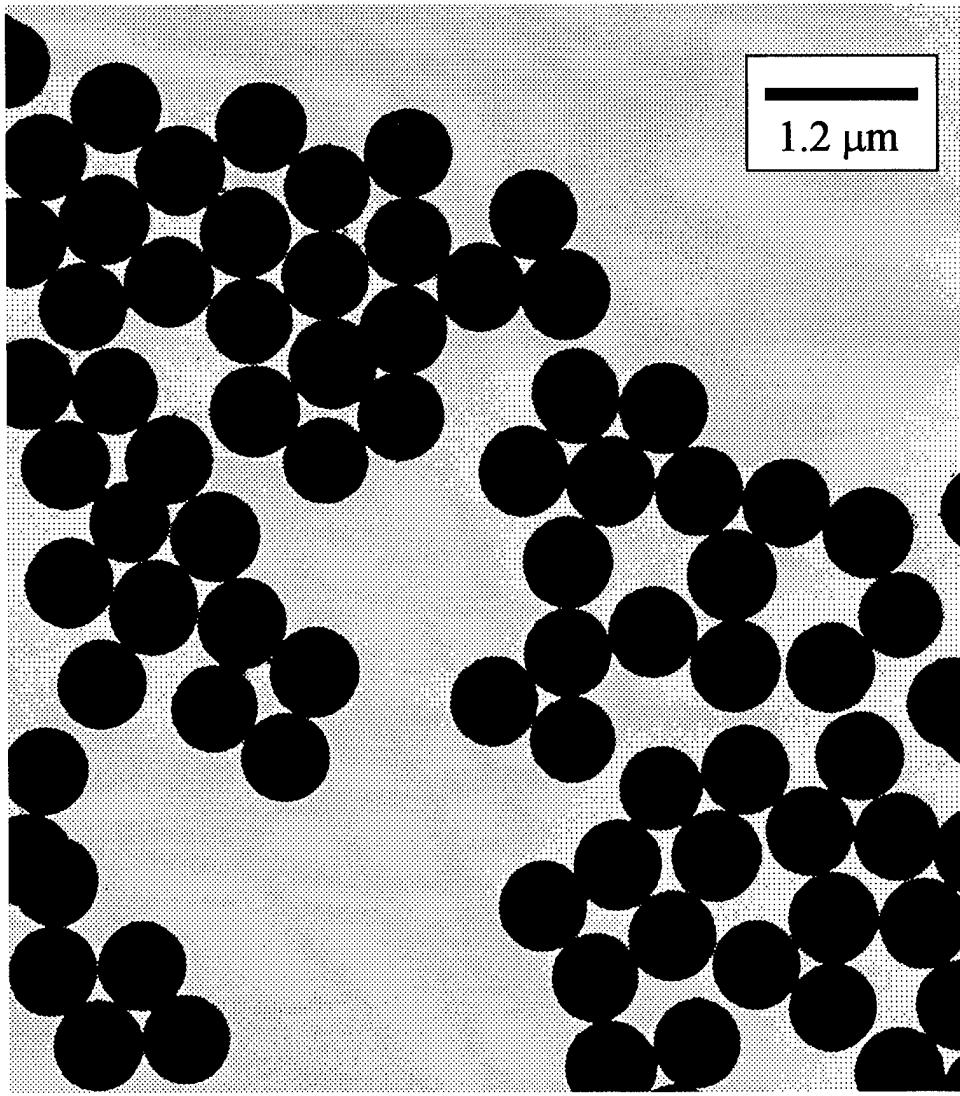


Figure 2. TEM Picture of Surface Modified SiO₂ Particles

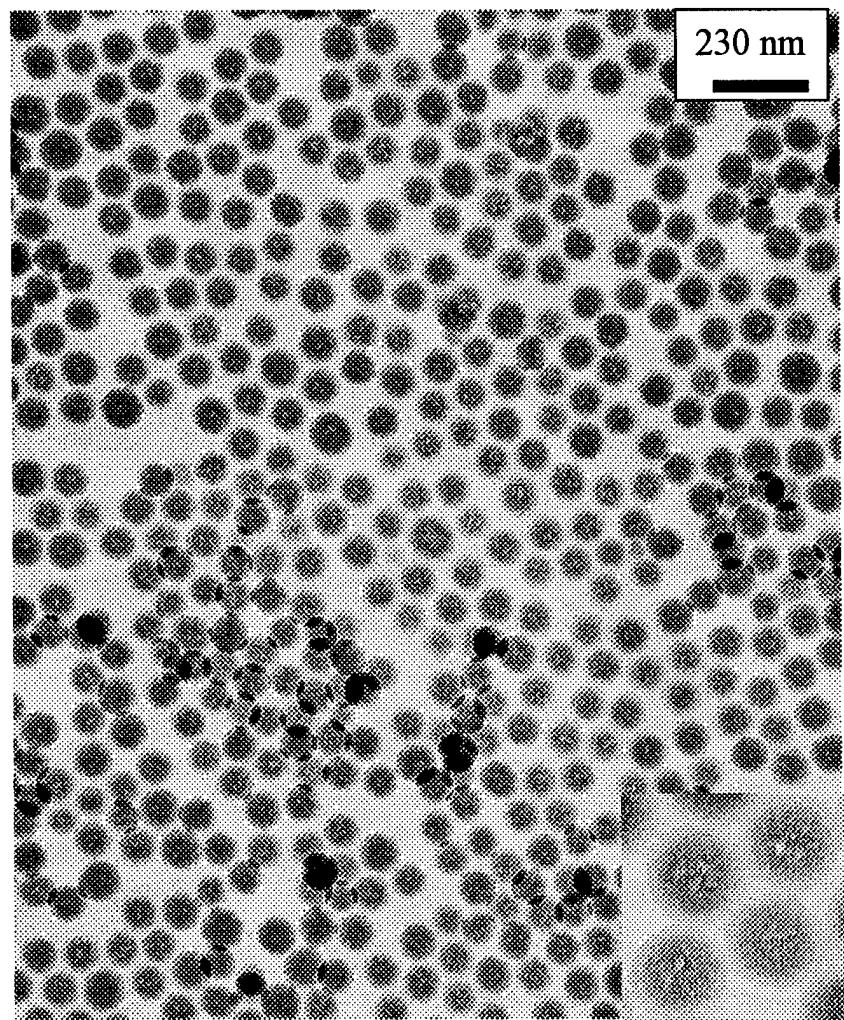


Figure 3. TEM picture of SiO₂-Ag composite particles
(Ag homogeneously distribution in SiO₂ spheres)

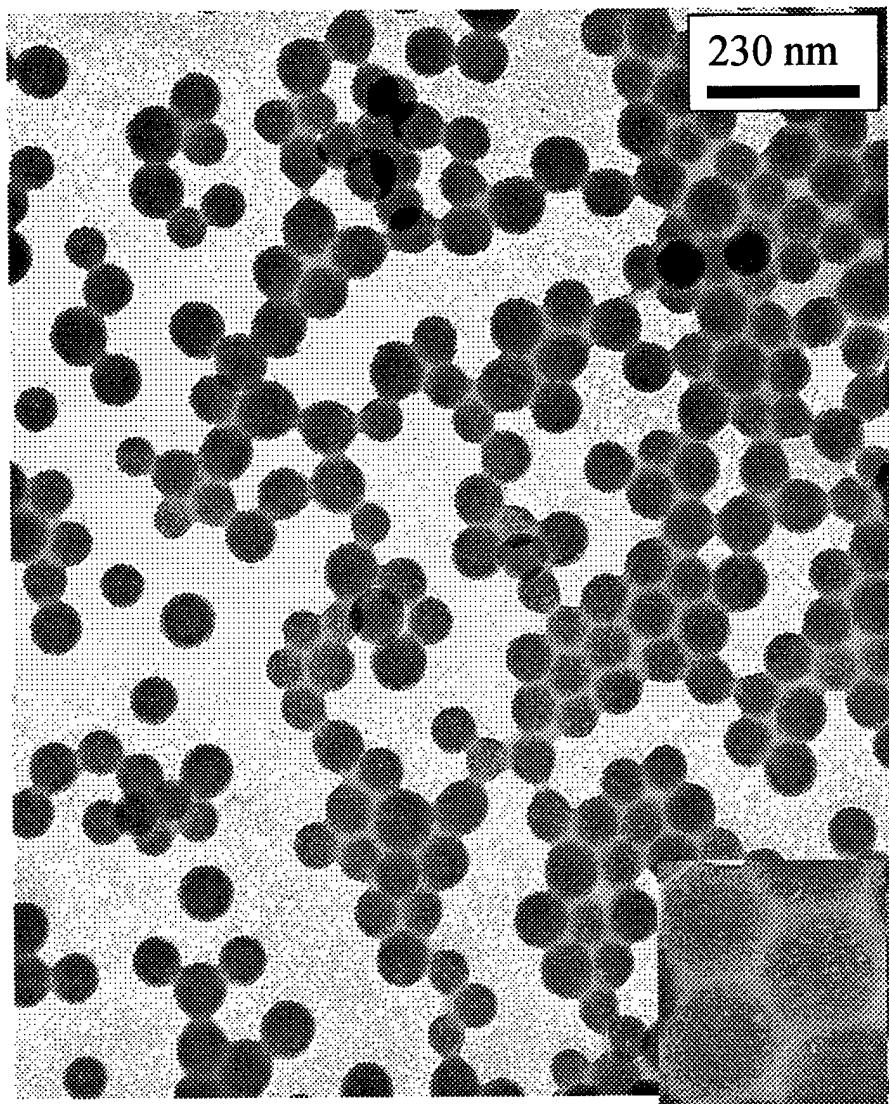


Figure 4. TEM picture of SiO_2 -Au composite spheres. The Au nanoparticles homogeneously distribute in the SiO_2 spheres.

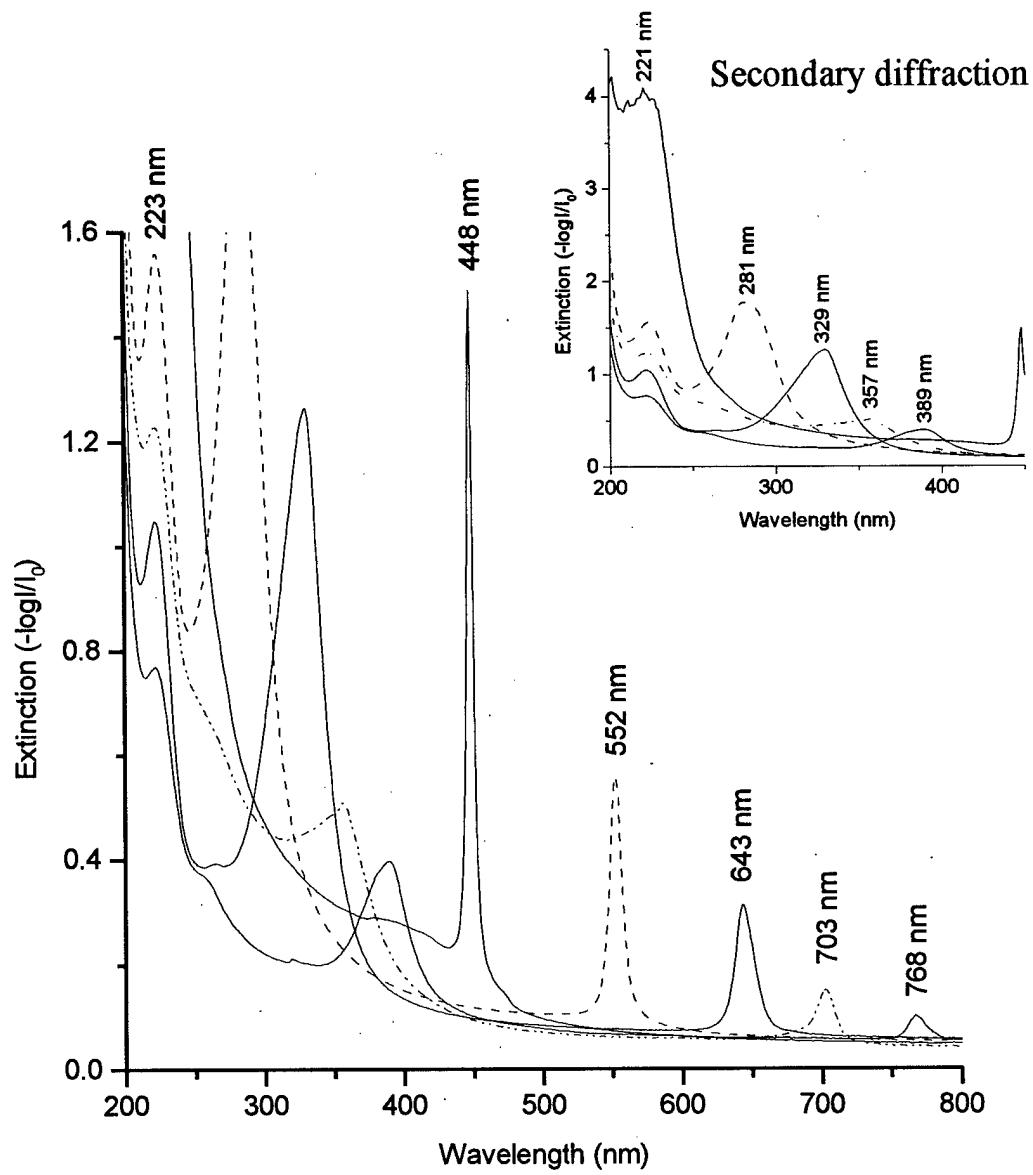


Figure 5. Extinction spectra of Ag@SiO₂ composite sphere CCA at different colloidal concentrations. The Ag@SiO₂ sphere diameter is 98±8 nm.

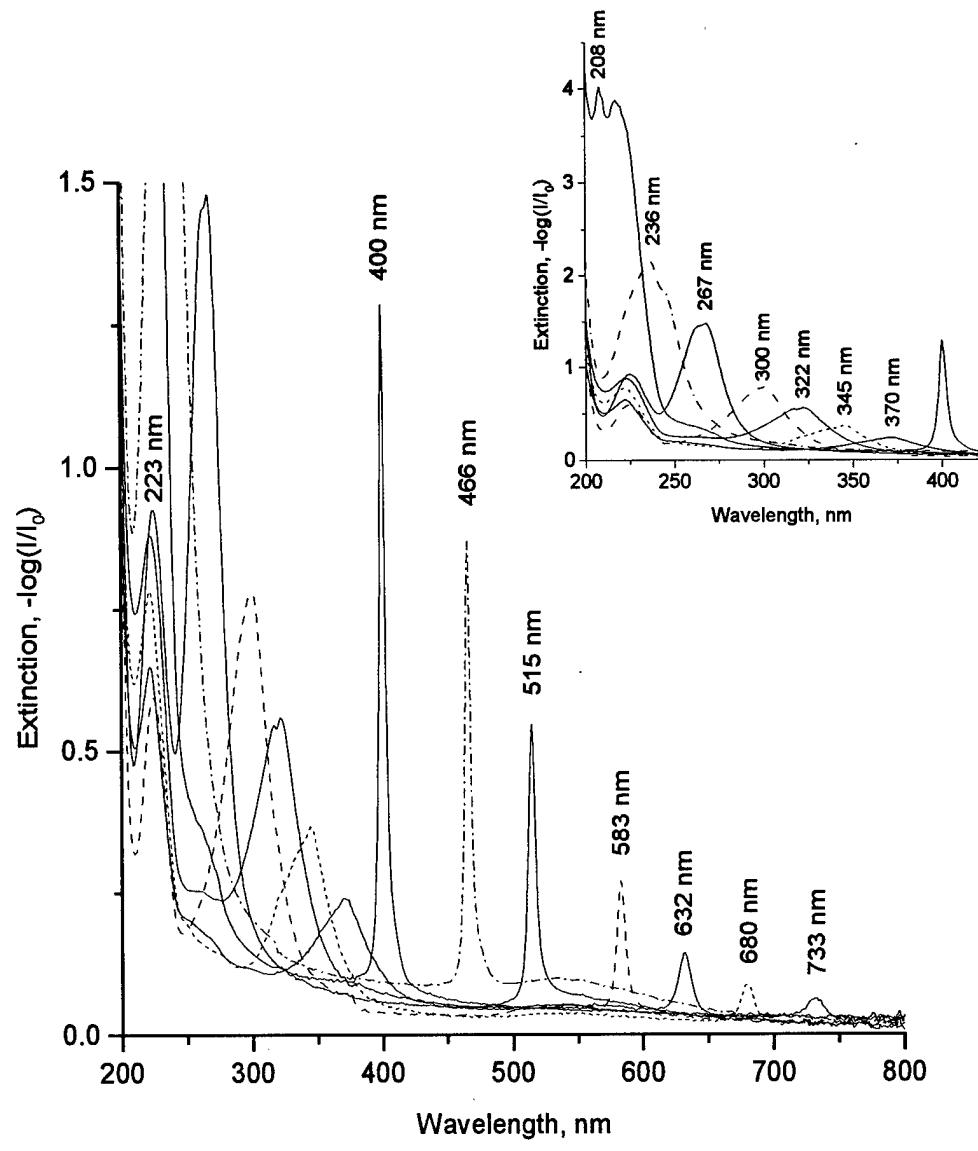


Figure 6. Extinction spectra of Au@SiO₂ composite sphere CCA at different sphere concentrations. The sphere diameter is 110±10 nm, and the thickness of the CCA is 0.125 mm.

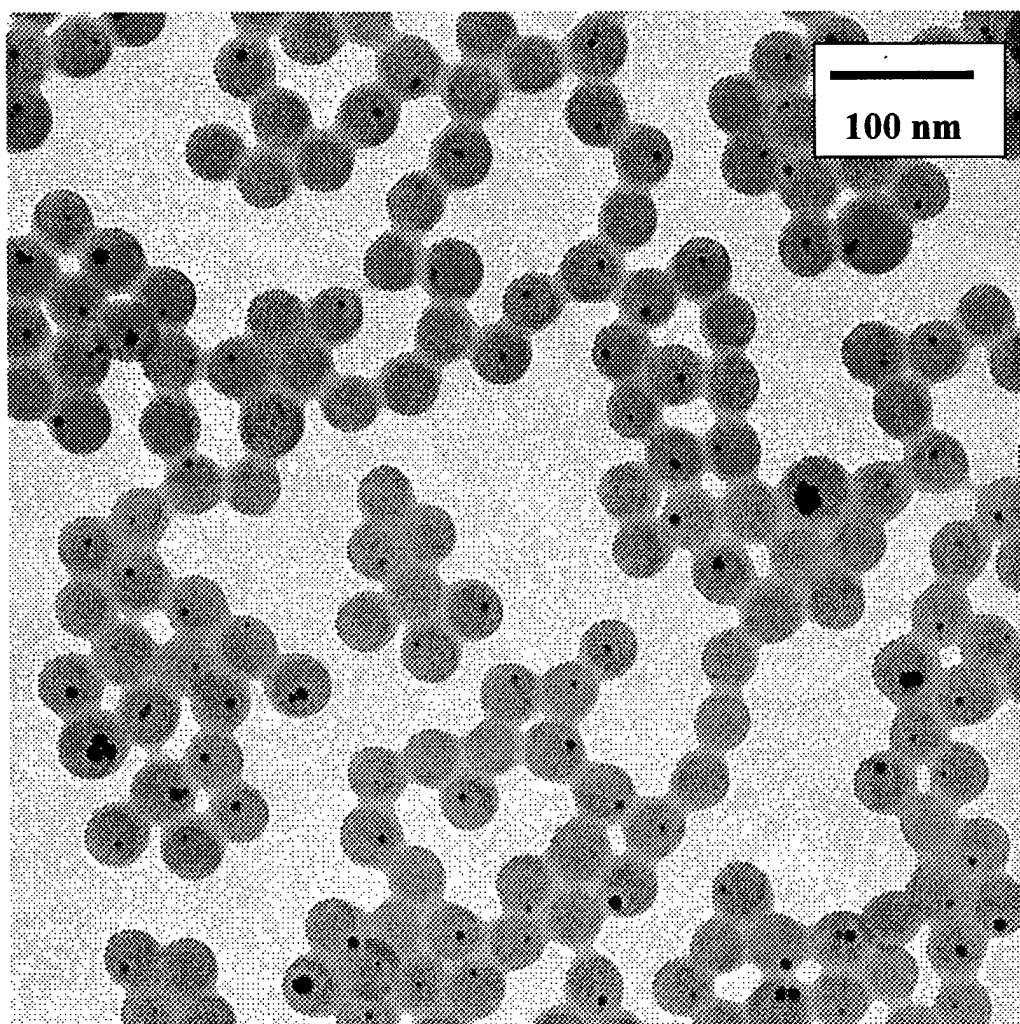


Figure 7. TEM picture of polystyrene-silver shell-core particles.

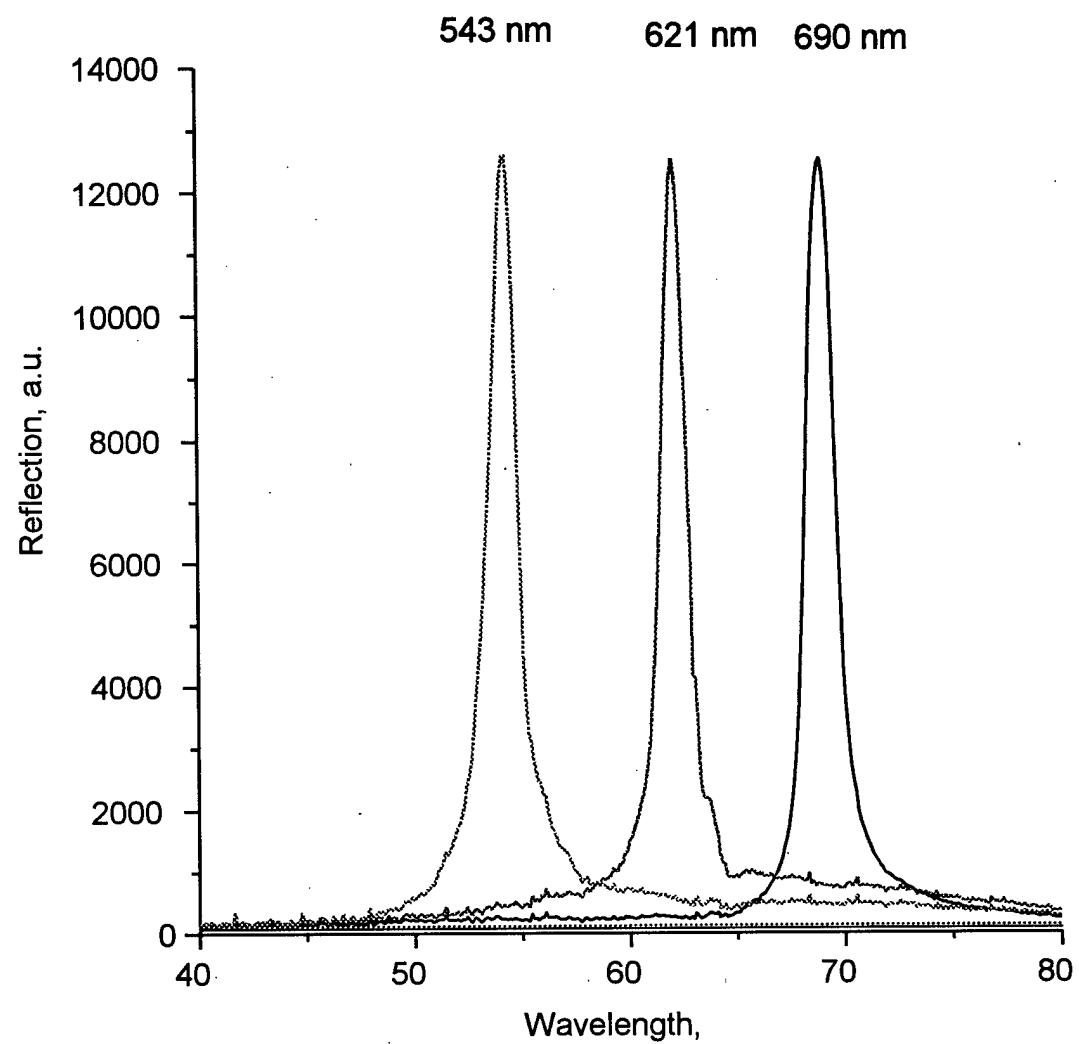


Figure 8. Back-scattering diffraction spectra of polystyrene-silver shell-core composite CCA at different particle concentrations.

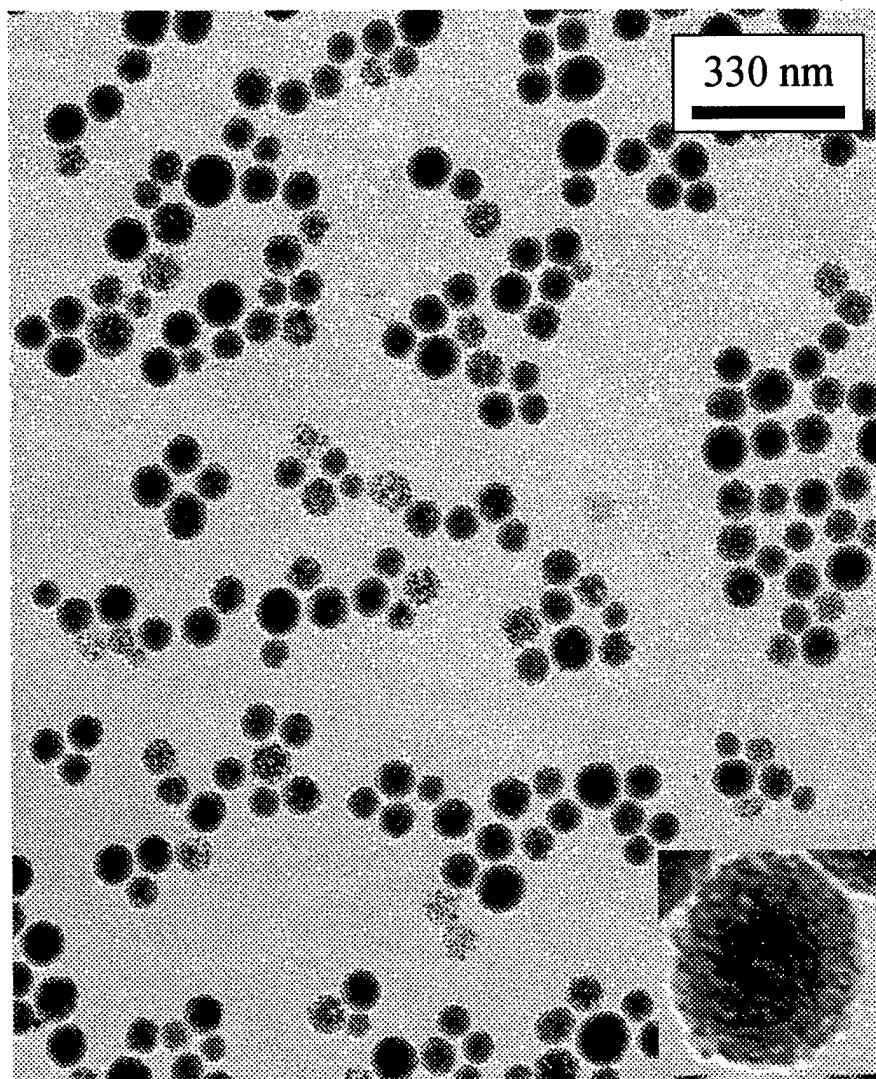


Figure 9. TEM picture of SiO₂-CdS (surface) composite particles

PCCA SWELLED INTO IR

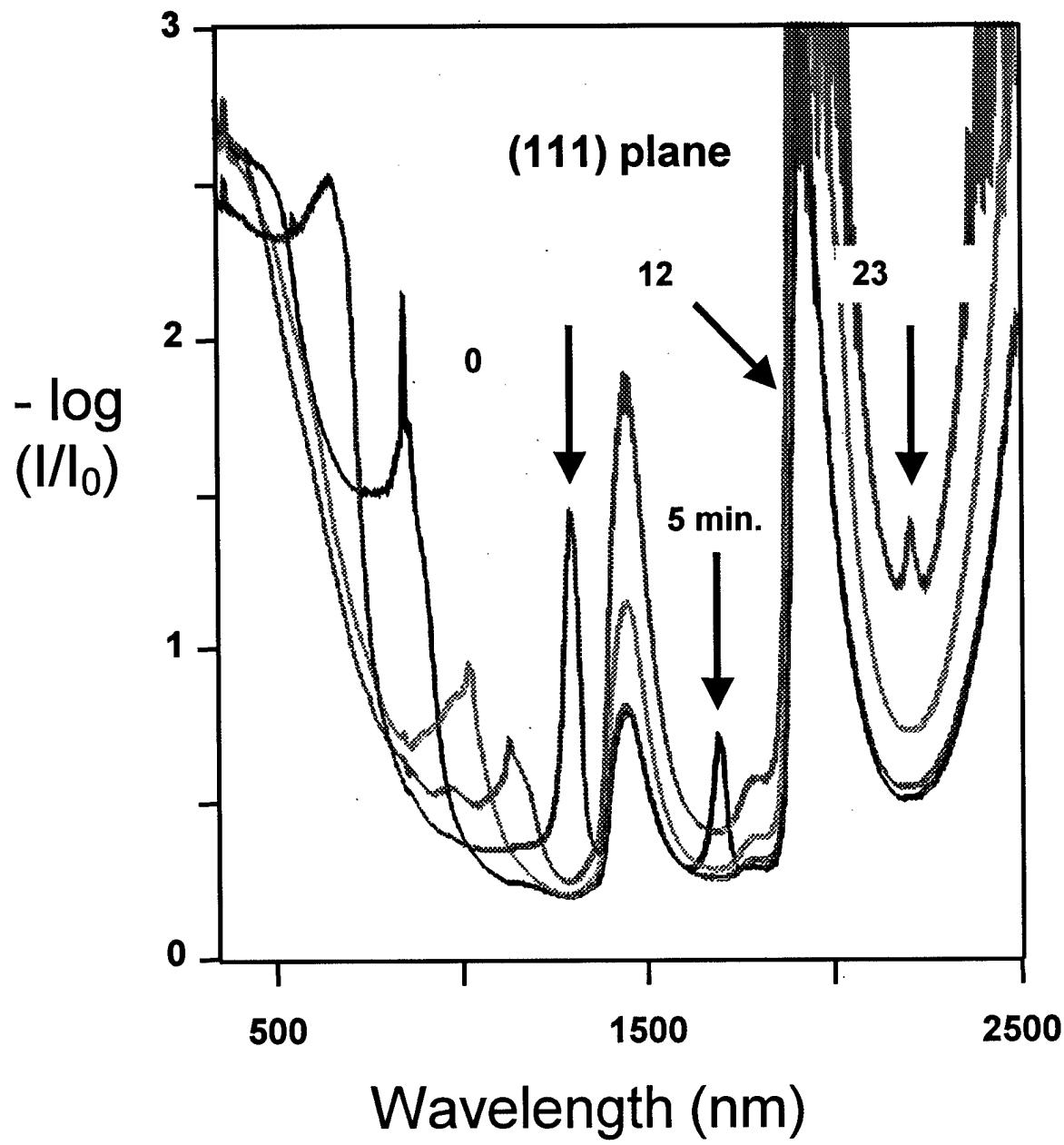


Figure 10. PS Colloid 270 nm diameter,
Acrylamide PCCA Hydrolyzed for 0, 5, 12, 23
minutes.

Increasing PCCA Spacing and Decreasing Elastic Constant Chemically Variable Crosslinking

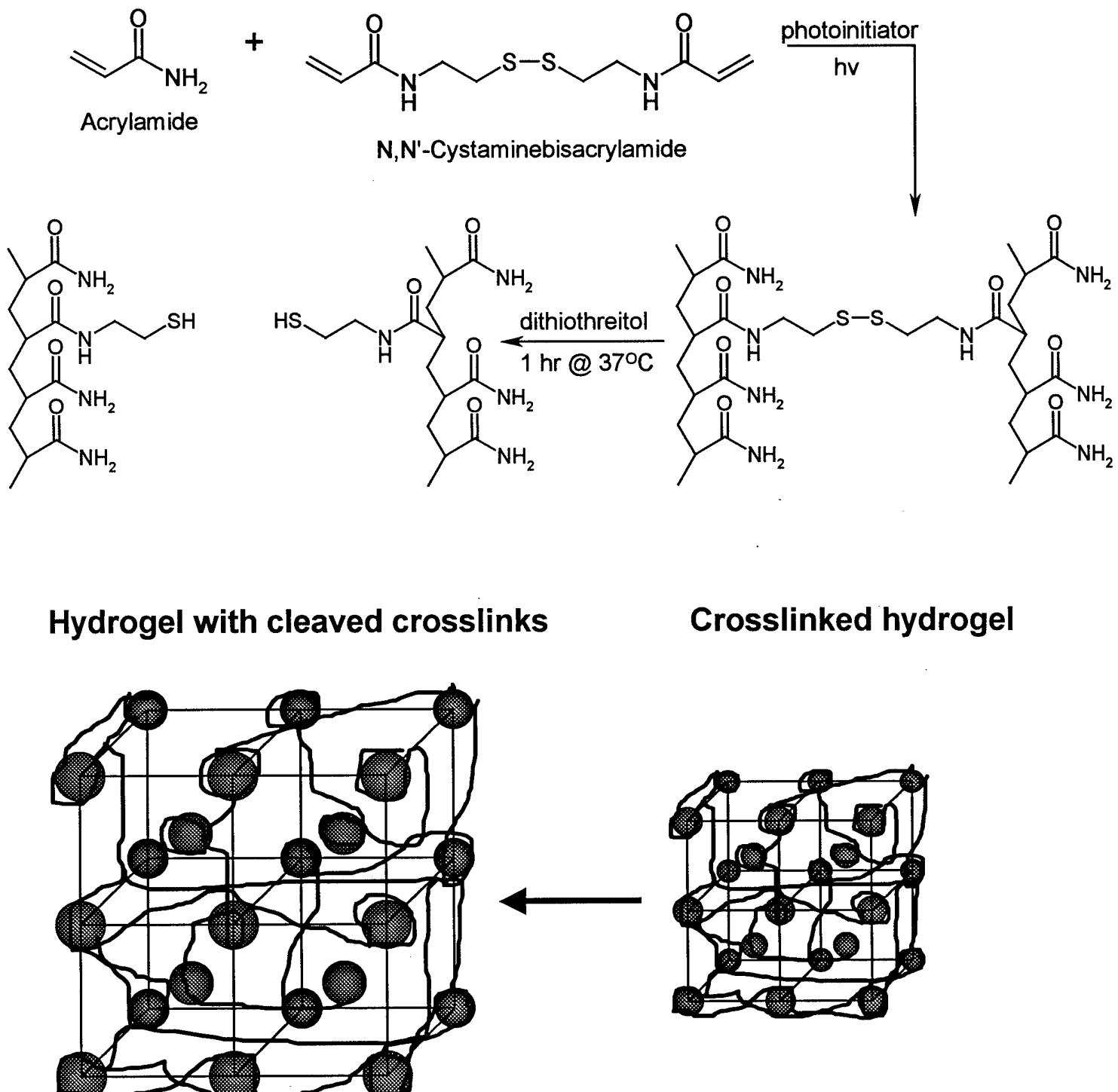


Figure 11

Fabrication of IR Diffracting Hydrogels

Synthesis of PEG Crosslinked Hydrogels

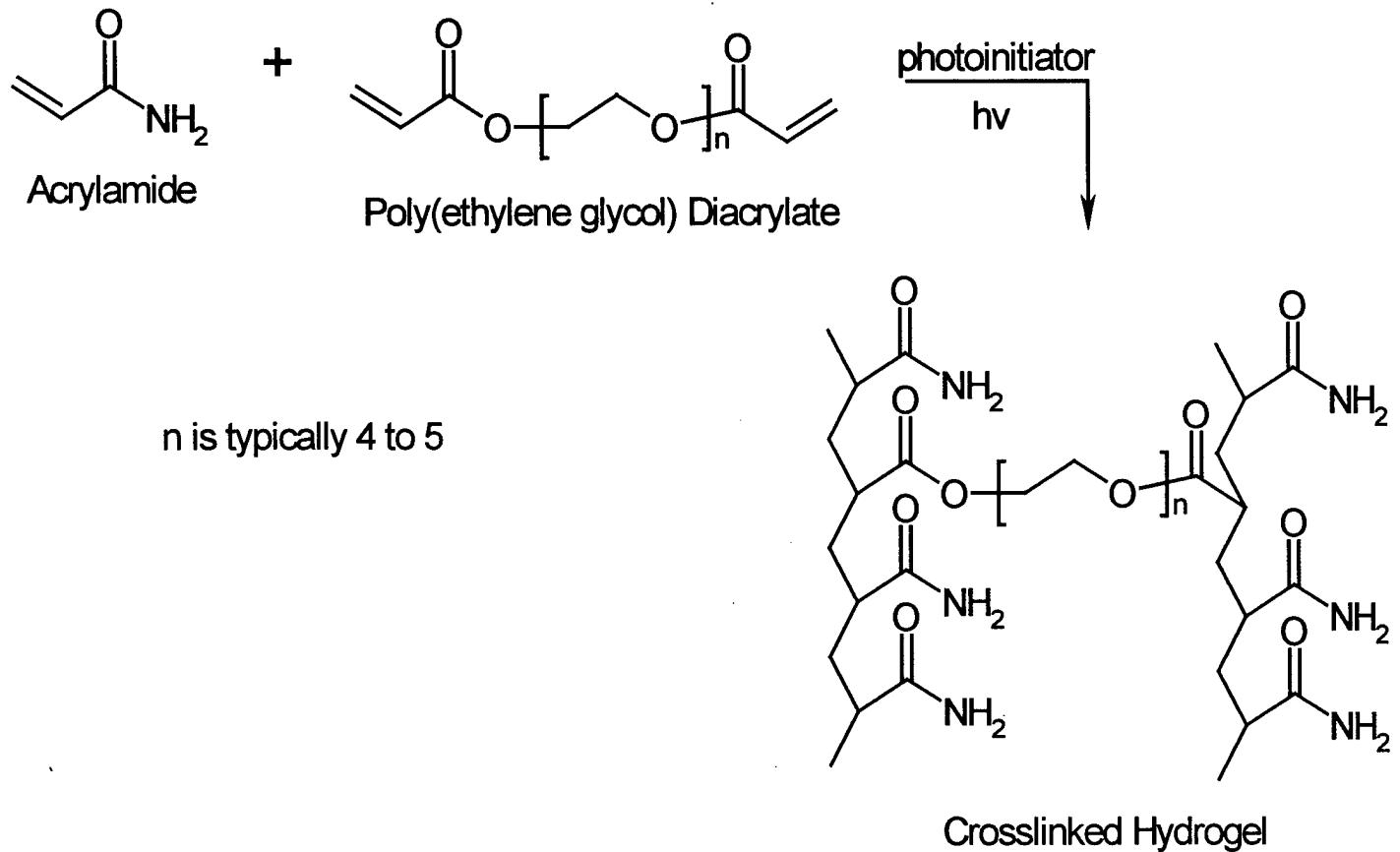
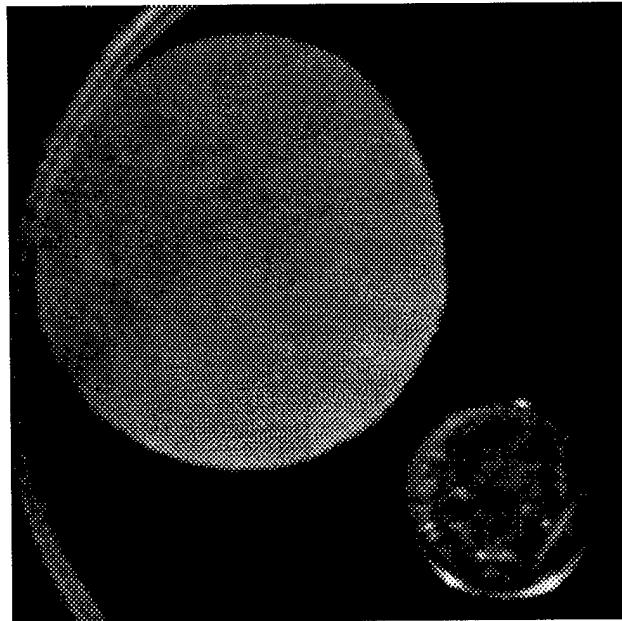
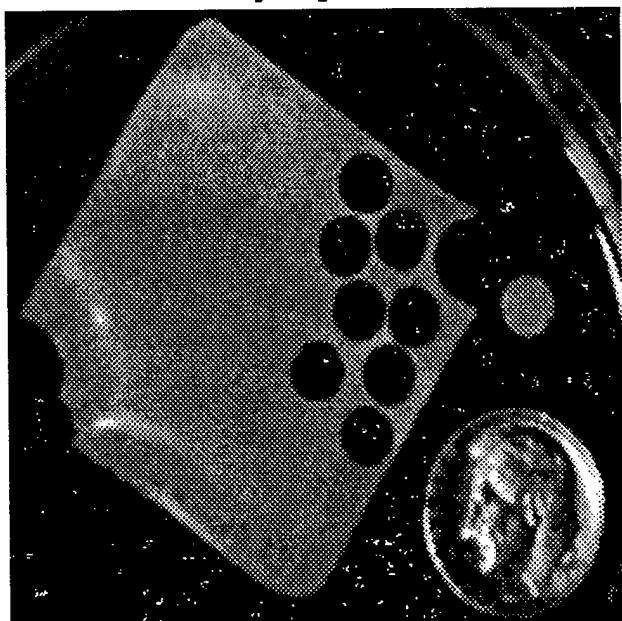
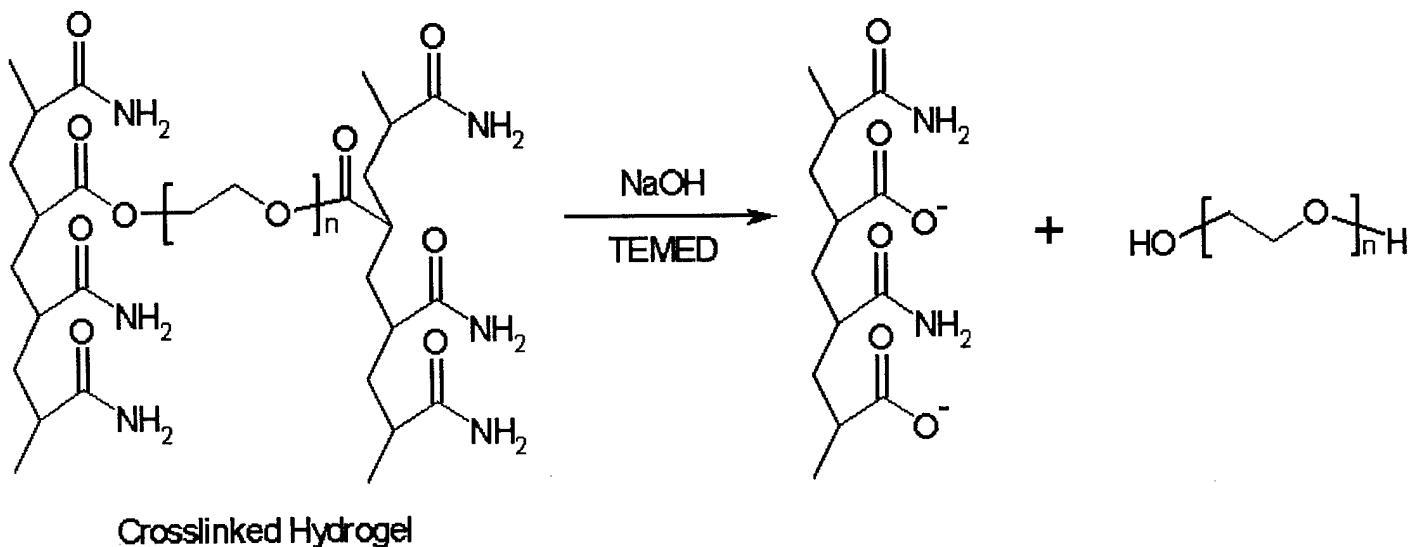


Figure 12

Fabrication of IR Diffracting Hydrogels

Hydrolysis of Crosslinks



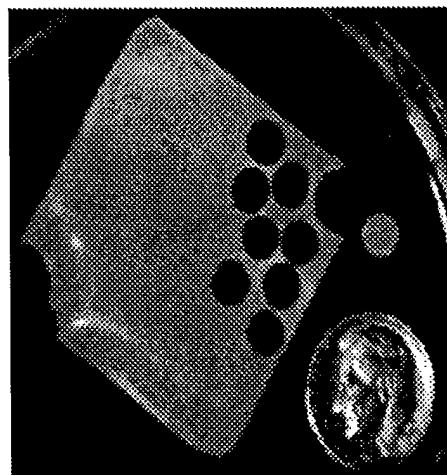
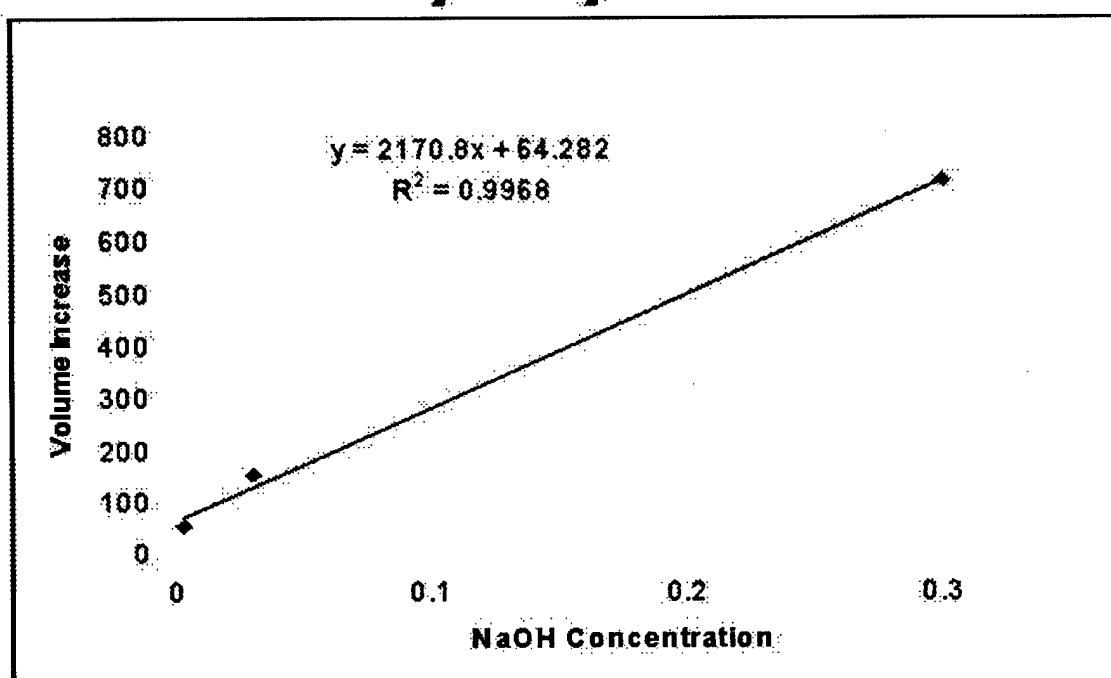
Typical Recipe:

0.03 M NaOH

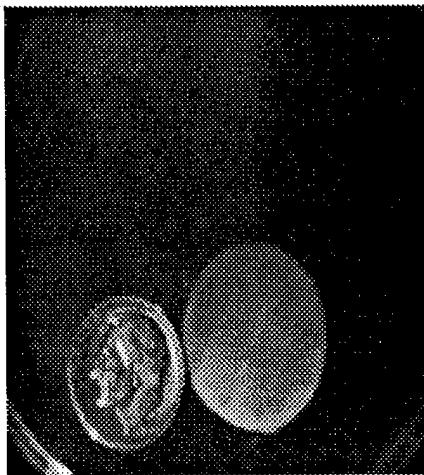
0.08% N,N,N',N'-tetramethylethylenediamine (TEMED)
for 1 hour

Figure 13

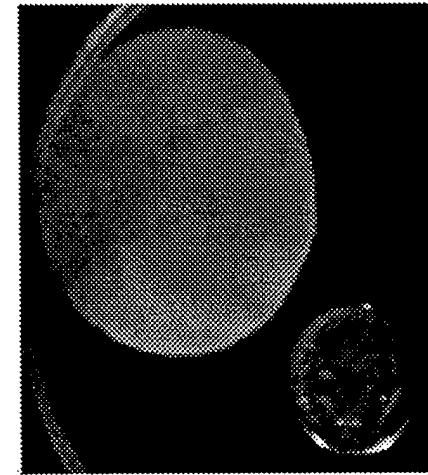
Gel Volume Increase With Changing NaOH Concentration After 1 Hour Of Hydrolysis



Gel Before Hydrolysis



After Hydrolysis
in 0.03 M NaOH



After Hydrolysis
in 0.3 M NaOH

$$\frac{V_f}{V_i} = 150$$

$$\frac{V_f}{V_i} = 10^3$$

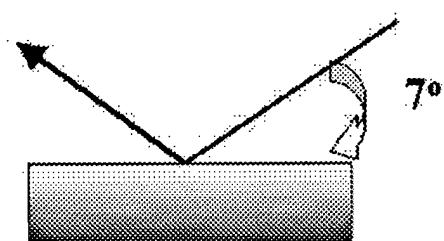
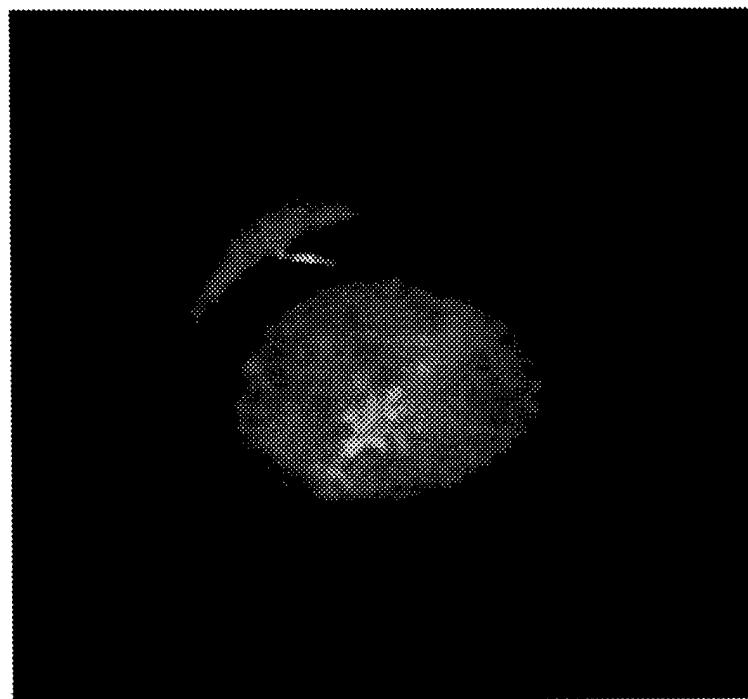
Figure 14

Diffraction Still Present After Swelling

$$\lambda_{\text{nn}}^i = 950 \text{ nm} \quad d_{\text{nn}}^i = 354 \text{ nm}$$

$$\frac{V_f}{V_i} = 150$$

$$\lambda_{\text{nn}}^f = 5.035 \mu\text{m} \text{ (Normal incidence)}$$



$$\lambda_d^r = 608 \text{ nm}$$

Figure 15

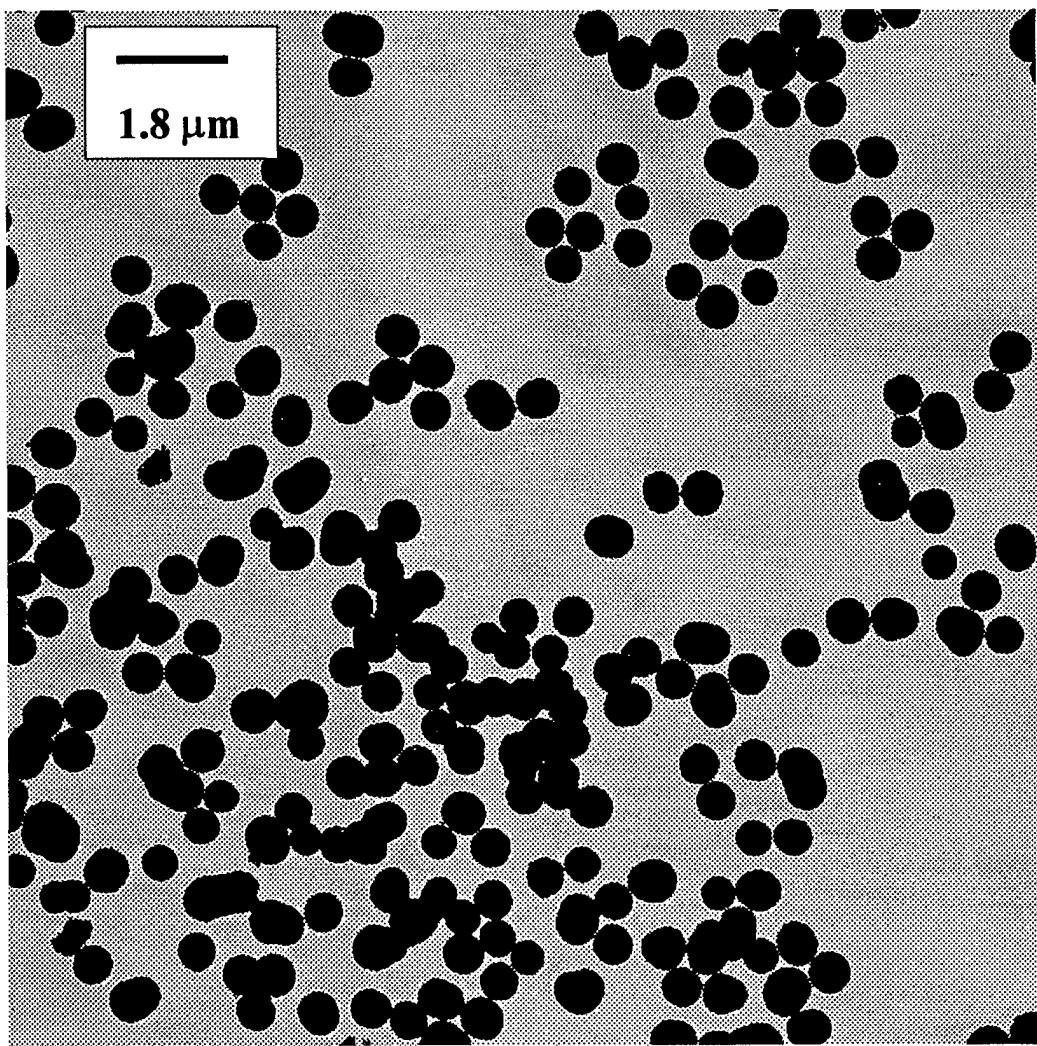


Figure 16. TEM picture of monodisperse TiO₂ spheres.